
(12) UK Patent Application (19) GB (11) 2 017 524 A

(21) Application No 7909712

(22) Date of filing 20 Mar 1979

(23) Claims filed 20 Mar 1979

(30) Priority data

(31) 7803075

(32) 22 Mar 1978

(33) Netherlands (NL)

(43) Application published
10 Oct 1979

(51) INT CL⁸

B01D 53/16

(52) Domestic classification
B1L 102 203 209 211 219

221 314 AE

C5E 181 PH

(56) Documents cited

GB 1393438

GB 1279637

GB 457343

(58) Field of search

B1L

C1A

C5E

(71) Applicants

Shell Internationale
Research Maatschappij
B.V.

Carel van Bylandtlaan 30,
The Hague, The
Netherlands

(72) Inventors

Roelof Cornelisse
Cornelis Ouwerkerk

(74) Agents

R C Rogers

(54) Removal of acid gases from a gas mixture

(57) Acid gases e.g. H₂S, CO₂, COS, are removed from a gas mixture by contacting at elevated pressure with an aqueous solution of a dialkyl - mono - alkanolamine (in particular diethyl - mono - ethanolamine), which aqueous solution does not contain a sterically hindered amine, i.e. a compound containing a secondary amine group bonded to a secondary or tertiary carbon atom, or a primary amine group bonded to a tertiary carbon atom. A physical solvent (e.g., sulfolane) may be present in the aqueous solution. The gas mixture may be natural gas, a gas obtained by the gasification of petroleum or coal, or a gas obtained by conversion of carbon monoxide and water into carbon dioxide and hydrogen.

GB 2 017 524 A

SPECIFICATION

A process for the removal of acid gases from a gas mixture

5

The invention relates to a process for the removal of acid gases from a gas mixture.

10

For many years aqueous solutions of alkanolamines have been used as absorbents for the removal of acid gases (by which are meant in this application hydrogen sulphide and carbon dioxide and also compounds which can readily be converted thereinto, such as carbon oxysulphide) from liquids, for example hydrocarbon mixtures and in particular 15 from gases which are at elevated pressure, such as natural gas, refinery gases. These solutions are suitable to reduce to lower values the acid gas contents of the gases or liquids to be purified.

20

The reduction of the content of acid gases, in particular H₂S, to a minimum is becoming increasingly important on account of demands made by the authorities in connection with environmental hygiene.

25

The gases to be purified are generally contacted with the absorbent at elevated pressure in an absorption column, with the result that the absorbent becomes loaded with acid gases. The regeneration of the loaded absorbent (i.e., the removal of all or the greater part of the absorbed gases therefrom) generally 30 occurs by heating it with steam in a regeneration column. Since steam (i.e., the energy to prepare steam) is expensive, the aim will be to keep the quantity of loaded absorbent as small as possible. It will be clear that on the one hand the aim is to use 35 absorbents having a high acid gas loading capacity and on the other hand, if one intends to remove in particular H₂S from CO₂-containing gas mixtures, to use absorbents which enable H₂S to be removed selectively with respect to CO₂ from gas mixtures.

40

The use of the smallest possible quantity of loaded absorbent to be regenerated is not only favourable to steam consumption, but the absorbent circulation system can also be constructed smaller than in the case of absorbents having a lower loading capacity 45 and/or selectivity.

45

A type of alkanolamines has now been found of which the aqueous solutions are selective to H₂S with respect to the absorption of H₂S from gas mixtures also containing CO₂ and also have a large 50 capacity for absorbing acid gases.

50

The invention therefore relates to a process for the removal of acid gases from a gas mixture, characterized in that said gas mixture is contacted at elevated pressure with an aqueous solution of a dialkyl - mono - alkanolamine, which aqueous solution does 55 not contain a sterically hindered amine.

55

By elevated pressure is meant a pressure of at least 2 bar. Preferably, the pressure is at least 5, in particular at least 10 bar.

60

By aqueous solution is meant a solution which contains at least 5% by weight of water.

65

A sterically hindered amine is defined as a compound which contains either a secondary amine group bound to a secondary or a tertiary carbon atom, or a primary amine group bound to a tertiary

carbon atom.

The two alkyl groups in the dialkyl - mono - alkanolamine need not be the same; methyl, ethyl, propyl and isopropyl groups are very suitable.

70 As very suitable alkanol groups may be mentioned the ethanol, propanol and isopropanol group.

Diethyl - mono - ethanolamine is preferred as dialkyl - mono - alkanolamine.

Generally, at least 10% by weight of the aqueous 75 solution consists of dialkyl - mono - alkanolamine; this percentage is preferably between 15 and 60.

If one intends to remove H₂S selectively from gas mixtures containing CO₂ it is advantageous to use an absorption column which contains a relatively small 80 number of trays, while the gas velocity will be chosen relatively high.

It is usual to contact the gas to be treated and the absorbent - in the present case the dialkyl - mono - alkanolamine solution - countercurrently with each

85 other in a tray column. The treated gas then leaves the vertical column near or at the top thereof, and the loaded absorbent, which contains the absorbed hydrogen sulphide and carbon dioxide, leaves said column near or at its bottom. Tray columns usually

90 contain valve trays, bubble cap trays or perforated sheets. Other types of trays or sheets may also be used. It is also possible to use packed columns.

The gas to be treated is preferably contacted with the absorbent at temperatures below 80°C, in particular 95 between 15 and 50°C.

The regeneration of the loaded aqueous solution of a dialkyl - mono - alkanolamine is suitably carried out in a regeneration column by heating the loaded absorbent and/or stripping it with an inert gas, such

100 as, for example, steam, preferably at temperatures of 80-120°C. Consequently, a gas containing H₂S, CO₂ and other components, if any, is separated from the loaded absorbent and a regenerated absorbent is obtained. The solution to be regenerated can very 105 suitably be heated by means of indirect heating with low-pressure steam. It is also possible to use direct injection of steam.

The aqueous solutions of dialkyl - mono - alkanolamines according to the invention have a large

110 capacity for the absorption of acid gases and are therefore also very suitable for the non-selective removal of CO₂ from gas mixtures at elevated pressure, since as a result of the relatively large quantity of CO₂ which can be absorbed, the quantity of absorbent to be used and consequently the quantity of

115 heat (usually steam) required for the regeneration of the loaded absorbent are smaller than when using amines with a lower capacity. For such removal of CO₂ it is of advantage that a physical solvent for acid gases is also present in the aqueous solution of the dialkyl - mono - alkanolamine, since as a result thereof the capacity for absorption and/or the rate of absorption of the absorbent is further increased,

120 foam formation is inhibited and any volatile mercaptans present in the gases to be purified are removed therefrom. Moreover, the steam consumption per volume unit of solvent to be regenerated is also favourably affected.

As physical solvents are very suitable N-methyl

130 pyrrolidone, propylene carbonate, methanol and in

particular tetrahydrothiophene dioxide (sulfolane). The quantity of physical solvent may vary between wide limits and may very suitably be 10-40% by weight of the aqueous solution.

5 The process according to the invention is very suitable to be used for the removal of CO₂ and/or H₂S and/or COS from natural gas, from gases obtained by the gasification of petroleum or coal, and from gases obtained by converting CO with H₂O into CO₂ and H₂.

EXAMPLE 1

A gas obtained in the gasification of coal (composition 37.1% by vol. of CO₂, 0.41% by vol. of H₂S, balance H₂) was treated countercurrently with an aqueous 2-molar solution of diethyl mono-ethanolamine (DEMEA) (according to the invention), and diisopropanolamine (DIPA) and methyl diethanolamine (MDEA) (both not according to the invention), respectively, in a tray column containing 55 trays at a pressure of 33 bar and a temperature of 40°C. The gas velocity was 20 cm/sec. The mutual ratio of the required volumes of the aqueous solutions of the amines in order to obtain a purified gas containing less than 4% by volume of CO₂ and less than 10 ppmv of H₂S was determined. It was found that if the volume required of the DEMEA-containing solution was assumed to be 1, the volume required of the MDEA- and DIPA- containing solutions was 1.59 and 2.86 respectively. In the process according to the invention the volume of aqueous amine solution to be used is therefore much smaller than in the known processes, so that the quantity of aqueous amine solution to be regenerated and recycled is much smaller than in known processes.

EXAMPLE 2

A gas mainly consisting of methane, which contained as contaminants 1.6% by vol. of CO₂ and 30 ppmv of H₂S, was treated countercurrently with absorbents containing 35 parts by weight of water, 15 parts by weight of sulfolane and 50 parts by weight of DEMEA, MDEA and DIPA, respectively, in an absorption column with 6 trays at a pressure of 80 bar and a gas velocity of 40 cm/sec. The mutual ratio of the required volumes of absorbent in order to obtain a gas having an H₂S-content of less than 3 ppmv of H₂S was determined, and the removed amount of CO₂ present in the starting gas was calculated. The table below shows that the required volume of the DEMEA-containing absorbent (according to the invention) is the smallest, while moreover the selectivity of this absorbent to H₂S removal in relation to CO₂ removal is higher than that of the DIPA-containing absorbent.

TABLE

Amine in absorbent	Required quantity of absorbent	CO ₂ absorbed in absorbent (% on quantity of CO ₂ in starting gas)
DEMEA	1	20.2
MDEA	1.43	20.1
DIPA	1.79	40.1

EXAMPLE 3

55 In order to purify a gas that mainly consisted of CH₄ and contained as contaminants 10% of CO₂, 8% of H₂S and 100 ppmv of CH₃SH, this gas was treated countercurrently (gas velocity 25 cm/sec.) with a solution containing 45 parts by weight of amine, 40 parts by weight of sulfolane and 15 parts by weight of water in a tray column with 30 trays at a pressure of 80 bar and a temperature of 40°C, at such a space velocity that the gas leaving the column contained less than 4 ppmv of H₂S and 5 ppmv of CH₃SH. The amines used were DEMEA and DIPA.

It was found that the required volume of the absorbent containing DIPA was 1.35 times larger than that of the absorbent containing DEMEA (according to the invention).

CLAIMS

1. A process for the removal of acid gases from a gas mixture, characterized in that said gas mixture is contacted at elevated pressure with an aqueous solution of a dialkyl - mono - alkanolamine, which aqueous solution does not contain a sterically hindered amine.
2. A process as claimed in claim 1, characterized in that the dialkyl - mono - alkanolamine used is diethyl - mono - ethanolamine.
3. A process as claimed in claims 1 and 2, characterized in that at least 10% by weight of the aqueous solution consists of dialkyl - mono - alkanolamine.
4. A process as claimed in claim 3, characterized in that said percentage is between 15 and 60.
5. A process as claimed in any one of the preceding claims, characterized in that the pressure is at least 5, preferably at least 10, bar.
6. A process as claimed in any one of the preceding claims, characterized in that the contacting temperature is between 15 and 50°C.
7. A process as claimed in any one of the preceding claims, characterized in that the aqueous solution of a dialkyl - mono - alkanolamine also contains a physical solvent for acid gases.
8. A process as claimed in claim 7, characterized in that the physical solvent used is tetrahydrothiophene dioxide.
9. A process as claimed in claim 7 or 8, characterized in that the quantity of physical solvent is 10-40% by weight of the aqueous solution.
10. A process as claimed in claim 1, substantially as described with special reference to the Examples.

Printed for Her Majesty's Stationery Office by The Tweeddale Press Ltd., Berwick-upon-Tweed, 1978.
Published at the Patent Office, 25 Southampton Buildings, London, WC2A 1AY, from which copies may be obtained.